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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/734,490	12/11/2003	Mark Zdeblick	021308-000510US	1638	
61487 7590 12/28/2006 BOZICEVIC, FIELD & FRANCIS LLP(PRTS) (PROTEUS BIOMEDICAL,INC) 1900 UNIVERSITY AVENUE, SUITE 200			EXAMINER		
			NASSER, ROBERT L		
	SITY AVENUE, SUITE 20 LTO, CA 94303		ART UNIT	PAPER NUMBER	
			3735		
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SHORTENED STATUTOR	Y PERIOD OF RESPONSE	MAIL DATE	DELIVER	DELIVERY MODE	
31 D	PAYS	12/28/2006 PAPER		PER	

Please find below and/or attached an Office communication concerning this application or proceeding.

If NO period for reply is specified above, the maximum statutory period will apply and will expire 6 MONTHS from the mailing date of this communication.

		Application No.	Applicant(s)				
Office Action Summary		10/734,490	ZDEBLICK ET AL.				
		Examiner	Art Unit				
· .		Robert L. Nasser	3735				
The MAILING DATE of Period for Reply	this communication app	pears on the cover sheet with the	correspondence address				
WHICHEVER IS LONGER, F - Extensions of time may be available ur after SIX (6) MONTHS from the mailing - If NO period for reply is specified above - Failure to reply within the set or extend	ROM THE MAILING D ider the provisions of 37 CFR 1.1 idea of this communication. e, the maximum statutory period ed period for reply will, by statute ian three months after the mailin.	Y IS SET TO EXPIRE 1 MONTH ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be will apply and will expire SIX (6) MONTHS from the application to become ABANDON grate of this communication, even if timely file.	ON. timely filed om the mailing date of this communication. NED (35 U.S.C. § 133).				
Status	,						
1) Responsive to commur	nication(s) filed on 25 A	uaust 2006					
2a) This action is FINAL .		action is non-final.					
<u> </u>							
·— · · ·	closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213.						
Disposition of Claims							
4) Claim(s) 1-3.5-52 and 5	4)⊠ Claim(s) <u>1-3,5-52 and 54-95</u> is/are pending in the application.						
	4a) Of the above claim(s) is/are withdrawn from consideration.						
5) Claim(s) is/are a							
	6) Claim(s) is/are rejected.						
		tion and/or election requiremen					
8) Claim(s) <u>1-3, 5-52, 54-95</u> are subject to restriction and/or election requirement. Application Papers							
•							
9) The specification is objected to by the Examiner.							
10) ☐ The drawing(s) filed on is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.							
		drawing(s) be held in abeyance. S					
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).							
11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.							
Priority under 35 U.S.C. § 119							
 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of: 1. Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received. 							
Attachment(s) 1) Notice of References Cited (PTO-8 2) Notice of Draftsperson's Patent Dra 3) Information Disclosure Statement(s Paper No(s)/Mail Date	awing Review (PTO-948)	4) Interview Summa Paper No(s)/Mail 5) Notice of Informal 6) Other:					

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Restriction to one of the following inventions is required under 35 U.S.C. 121:

- Claims 1-47, drawn to a multiplexed medical carrier, classified in class 606, subclass 1.
- II. Claims 48-68, drawn to a method for configuring a medical carrier, classified in class 600, subclass 486.
- III. Claims 69-82, drawn to a method of collecting medical data, classified in class 600, subclass 300.
- IV. Claims 83-95, drawn to a method of delivering energy or one or more substances to the body, classified in class 607, subclass 1.

The inventions are distinct, each from the other because of the following reasons:

Inventions I and II are related as process of making and product made. The inventions are distinct if either or both of the following can be shown: (1) that the process as claimed can be used to make another and materially different product or (2) that the product as claimed can be made by another and materially different process (MPEP § 806.05(f)). In the instant case, the product could be made by a materially different method, such as one where the conductor runs on the exterior of a substrate with no surface penetration.

Inventions I and III are related as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product. See MPEP § 806.05(h). In the instant case, the product

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can be used to practice a materially different method, such as providing energy to tissue.

Inventions I and IV are related as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product. See MPEP § 806.05(h). In the instant case, the product can be used to practice a materially different method, such as monitoring blood pressure.

Inventions II and III are directed to related processes. The related inventions are distinct if the (1) the inventions as claimed are either not capable of use together or can have a materially different design, mode of operation, function, or effect; (2) the inventions do not overlap in scope, i.e., are mutually exclusive; and (3) the inventions as claimed are not obvious variants. See MPEP § 806.05(j). In the instant case, the inventions as claimed are not capable of use together, as a method of configuring a device is not useable with a method of using the device. Furthermore, the inventions as claimed do not encompass overlapping subject matter and there is nothing of record to show them to be obvious variants.

Inventions II and IV are directed to related processes. The related inventions are distinct if the (1) the inventions as claimed are either not capable of use together or can have a materially different design, mode of operation, function, or effect; (2) the inventions do not overlap in scope, i.e., are mutually exclusive; and (3) the inventions as

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claimed are not obvious variants. See MPEP § 806.05(j). In the instant case, the inventions as claimed are not capable of use together, as a method of configuring a device is not useable with a method of using the device. Furthermore, the inventions as claimed do not encompass overlapping subject matter and there is nothing of record to show them to be obvious variants.

Inventions III and IV are directed to related processes. The related inventions are distinct if the (1) the inventions as claimed are either not capable of use together or can have a materially different design, mode of operation, function, or effect; (2) the inventions do not overlap in scope, i.e., are mutually exclusive; and (3) the inventions as claimed are not obvious variants. See MPEP § 806.05(j). In the instant case, the inventions as claimed operate in materially different mode of operation. Furthermore, the inventions as claimed do not encompass overlapping subject matter and there is nothing of record to show them to be obvious variants.

Because these inventions are independent or distinct for the reasons given above and there would be a serious burden on the examiner if restriction is not required because the inventions have acquired a separate status in the art due to their recognized divergent subject matter, restriction for examination purposes as indicated is proper.

If applicant elects groups I or II, the following election of species is also required.

Species I, where the transducer is a sensor.

Species II, where the transducer is an actuator.

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These are distinct since they operate in different modes of operation.

If applicant elects group III or IV, then the following elections of species requirement is deemed pertinent.

Species I, where the data is collected from a catheter present in one or more blood vessels or a heart chamber.

Species II, where the data is collected from a flat surface present on or near brain tissue.

Species III, where the data is collected from a catheter present in a urinary tract.

Species IV, where the data is collected from a catheter present in reproductive tract.

Species V, where the data is collected from a catheter present in an endoscopic surgical site.

Species VI, where the data is collected from a catheter present in an abdominal cavity.

Species VII, where the data is collected from a catheter present in a gastrointestinal tract.

Species VIII, where the data is collected from a catheter present adjacent a bone or in a joint space.

The species are independent or distinct because they each sense data or supply energy in a materially different manner.

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Applicant is required under 35 U.S.C. 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable. Currently, claims 40-47 or no claims are generic.

Applicant is advised that a reply to this requirement must include an identification of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which depend from or otherwise require all the limitations of an allowable generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species.

MPEP § 809.02(a).

Applicant is advised that the reply to this requirement to be complete must include (i) an election of a species or invention to be examined even though the requirement be traversed (37 CFR 1.143) and (ii) identification of the claims encompassing the elected invention.

The election of an invention or species may be made with or without traverse. To reserve a right to petition, the election must be made with traverse. If the reply does not distinctly and specifically point out supposed errors in the restriction requirement, the election shall be treated as an election without traverse.

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Should applicant traverse on the ground that the inventions or species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the inventions or species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C.103(a) of the other invention.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Robert L. Nasser whose telephone number is 571 272-4731. The examiner can normally be reached on m-f 9-5.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Charles Marmor II can be reached on 571 272-4730. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

Robert L. Nasser Primary Examiner Art Unit 3735

RLN December 20, 2006

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